Serial No.: 09/904,356 Filed : July 12, 2001

Page 3

Amendments to the Claims

This listing of claims will replace all prior versions and listings of the claims in this application.

- --7. (Currently amended) A method of inhibiting fusion of a macrophage-tropic primary isolate of HIV-1 to a CD4+ cell susceptible to infection by a macrophage-tropic primary isolate of HIV-1 which comprises contacting the CD4+ cell with an effective amount of an agent which is (1) capable of inhibiting fusion of HeLa-env_{JR-FL} to a PM1 cell a macrophage-tropic primary isolate of HIV-1 to a CD4+ cell susceptible to infection by a macrophage-tropic primary isolate of HIV-1, but (2) not capable of inhibiting fusion of HeLa-env_{LAI} to a HeLa-CD4+ cell a T cell-tropic isolate of HIV-1 to a CD4+ cell susceptible to infection by a T cell-tropic isolate of HIV-1, so as to thereby inhibiting inhibit the fusion of the macrophage-tropic primary isolate of HIV-1 to the CD4+ cell.--
- --8. (Currently amended) The method of claim 7, wherein the agent is determined to be capable of inhibiting fusion of a macrophage-tropic primary isolate of HIV-1 to a CD4+ cell but not capable of inhibiting fusion of a T cell tropic isolate of HIV-1 to a CD4+ cell using a method which comprises:
- (a) contacting (i) a first appropriate CD4+ PM1 cell, which is labeled with a first dye, with (ii) a HeLa-env_{JR-FL} cell expressing an HIV-1 envelope glycoprotein of the macrophage-tropic primary isolate of HIV-1 on its surface, which is labeled with a second dye, in the presence of an excess of the agent under conditions which would normally permit the fusion of the CD4+ PM1 cell to the HeLa-env_{JR-FL} cell expressing the HIV-1 envelope glycoprotein on its surface in the absence of the agent, the first and second dyes being selected so as to allow resonance energy transfer between the dyes;

Serial No.: 09/904,356 Filed : July 12, 2001

Page 4

- (b) exposing the product of step (a) to conditions which would result in resonance energy transfer if fusion has occurred; and
- (c) determining whether there is a reduction of resonance energy transfer, when compared with the resonance energy transfer in the absence of the agent;
- (d) contacting (i) a second appropriate CD4+ HeLa-CD4+ cell, which is labeled with a first dye, with (ii) HeLa-env_{LAI} a cell expressing an HIV-1 envelope glycoprotein of a T cell-tropic isolate of HIV-1 on its surface, which is labeled with a second dye, in the presence of an excess of the agent under conditions which would normally permit the fusion of the CD4+ cell HeLa-CD4+ to the HeLa-env_{LAI} cell expressing the HIV-1 envelope glycoprotein on its surface in the absence of the agent, the first and second dyes being selected so as to allow resonance energy transfer between the dyes;
- (e) exposing the product of step (d) to conditions that would result in resonance energy transfer if fusion has occurred;
- (f) determining whether there is a reduction of resonance energy transfer, when compared with the resonance energy transfer in the absence of the agent; and
- (g) comparing the determination made in step (c) with the determination made in step (f), wherein a decrease in transfer in step (c) but not in step (f) indicates that the agent is capable of specifically inhibiting fusion of the macrophage-tropic primary isolate of HIV-1 to the CD4+ cell, but not capable of specifically inhibiting the fusion of a T cell-tropic isolate of HIV-1 to the CD4+ cell.—
- --9. (Previously presented) The method of claim 7, wherein the agent is an antibody.--

Claims 10-12 (Cancelled).

Serial No.: 09/904,356 Filed : July 12, 2001

Page 5

--13. (New) The method of claim 7, wherein the agent is capable of inhibiting fusion of a macrophage-tropic primary isolate of HIV-1 to a CD4+ cell but not capable of inhibiting fusion of a T cell-tropic isolate of HIV-1 to a CD4+ cell in a method which comprises:

- (a) contacting (i) a PM1 cell, which is labeled with a first dye, with (ii) $\text{HeLa-env}_{\text{JR-FL}}$, which is labeled with a second dye, in the presence of an excess of the agent under conditions which would normally permit the fusion of the PM1 cell to the HeLa-env_{JR-FL} in the absence of the agent, the first and second dyes being selected so as to allow resonance energy transfer between the dyes;
- (b) exposing the product of step (a) to conditions which would result in resonance energy transfer if fusion has occurred; and
- (c) determining whether there is a reduction of resonance energy transfer, when compared with the resonance energy transfer in the absence of the agent;
- (d) contacting (i) a HeLa-CD4+ cell, which is labeled with a first dye, with (ii) HeLa-env_{LAI}, which is labeled with a second dye, in the presence of an excess of the agent under conditions which would normally permit the fusion of HeLa-CD4+ to the HeLa-env_{LAI} in the absence of the agent, the first and second dyes being selected so as to allow resonance energy transfer between the dyes;
- (e) exposing the product of step (d) to conditions that would result in resonance energy transfer if fusion has occurred;
- (f) determining whether there is a reduction in resonance energy transfer, when compared with the resonance energy transfer in the absence of the agent; and
- (g) comparing the determination made in step (c) with the determination made in step (f), wherein a decrease in transfer in

Serial No.: 09/904,356 Filed : July 12, 2001

Page 6

step (c) but not in step (f) indicates that the agent is capable of specifically inhibiting fusion of the macrophage-tropic primary isolate of HIV-1 to the CD4+ cell, but not capable of specifically inhibiting the fusion of a T cell-tropic isolate of HIV-1 to the CD4+ cell.

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